



EC DECLARATION OF CONFORMITY

In Vitro Diagnostic Medical Devices for Professional use

DECLARATION N°: 04-06-04-01

We

Fuller Laboratories

1135 East Truslow Avenue Fullerton, CA 92831 USA
Telephone: 1 714 525 7660 Fax: 1 714 525 7614

having designated :

Medimark Europe Sarl ,

11 rue Emile Zola – 38033 Grenoble Cedex 2 - France

As our European Authorized Representative

insure and declare under our sole responsibility that the In Vitro Diagnostic Medical Devices specified in the attached list to which this declaration relates are in conformity with the applicable requirements of the In Vitro Diagnostic Medical Devices Directive 98/79/EC.

This declaration is made in accordance with Annex III of the In Vitro Diagnostic Medical Devices Directive 98/79/EC and is valid for an undetermined period of time .

Fullerton, 6 April 2004

A handwritten signature in black ink that reads "Lee Fuller". The signature is written in a cursive style with a long horizontal stroke at the end.

Lee Fuller, President



ANNEX TO DECLARATION N°: 04-06-04-01

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List of In Vitro Diagnostic Medical Devices in relation with the above declaration

Device designation / Ref N°	EDMS Class N°
Babesia microti IFA IgG Antibody Kit / BMG-120	14 05 02 90 00

Fullerton, 6 April 2004

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Lee Fuller, President